

Hypnosis and self-hypnosis, administered and taught by nurses, for the reduction of chronic pain: a controlled clinical trial

Submission date 06/09/2005	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input checked="" type="checkbox"/> Protocol
Registration date 04/10/2005	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 10/09/2009	Condition category Signs and Symptoms	<input type="checkbox"/> Individual participant data <input type="checkbox"/> Record updated in last year

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Dr Bernard Burnand

Contact details
IUMSP
Bugnon 17
Lausanne
Switzerland
CH-1010

Additional identifiers

Study information

Scientific Title

Study objectives
Hypnosis and auto-hypnosis do not reduce pain in patients with chronic pain

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Randomised controlled trial

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Various conditions engendering chronic pain

Interventions

Hypnosis and auto-hypnosis administered and taught by trained nurses

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

Level of pain (VAS)

Key secondary outcome(s)

1. Physical activity (tridimensional accelerometer)
2. Health related quality of life (SF-36)

Completion date

31/12/1994

Eligibility

Key inclusion criteria

1. To be referred (including occasional self-referral) for the management of chronic pain
2. To live within manageable distances from the hospital
3. To be aged 18 or above
4. To understand and speak French adequately
5. To be able to see, hear and communicate (i.e. no alteration of consciousness)
6. To have no diagnosis of mental illness (e.g. psychosis)
7. To give informed consent

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Lower age limit

18 years

Sex

All

Key exclusion criteria

Patients were excluded from the study whenever they wished and/or when no conventional treatment could be proposed.

Date of first enrolment

01/05/1993

Date of final enrolment

31/12/1994

Locations**Countries of recruitment**

Switzerland

Study participating centre

IUMSP

Lausanne

Switzerland

CH-1010

Sponsor information**Organisation**

Hospital of Morges (Switzerland)

ROR

<https://ror.org/026qe1m81>

Funder(s)

Funder type

Government

Funder Name

Swiss National Science Foundation (Switzerland) (SNF 4034-35883)

Alternative Name(s)

Schweizerischer Nationalfonds, Swiss National Science Foundation, Fonds National Suisse de la Recherche Scientifique, Fondo Nazionale Svizzero per la Ricerca Scientifica, Fonds National Suisse, Fondo Nazionale Svizzero, Schweizerische Nationalfonds, The Swiss National Science Foundation (SNSF), SNF, SNSF, FNS

Funding Body Type

Private sector organisation

Funding Body Subtype

Trusts, charities, foundations (both public and private)

Location

Switzerland

Results and Publications

Individual participant data (IPD) sharing plan**IPD sharing plan summary****Study outputs**

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Protocol article	protocol	01/08/1994		Yes	No